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4. The method of claim 1, wherein the subject has a fasting baseline LDL-C from about 40 mg/dl to about 115 mg/dl.

5. The method of claim 1, wherein subject has one or more of: a median baseline fasting non-HDL-C of about 200 mg/dl to about 300 mg/dl, a median baseline fasting total cholesterol of about 250 mg/dl to about 300 mg/dl, a median baseline fasting VLDL-C of about 140 mg/dl to about 200 mg/dl, and/or a median baseline fasting HDL-C of about 10 mg/dl to about 80 mg/dl.

6. The method of claim 1, comprising administering to the subject about 4 g of the pharmaceutical composition daily for the period of at least about 12 weeks to effect a reduction in fasting triglycerides of at least about 10% without substantially increasing LDL-C compared to placebo control.

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7. The method of claim 1, comprising administering to the subject about 4 g of the pharmaceutical composition daily for the period of at least about 12 weeks to effect a reduction in fasting triglycerides of at least about 25% without substantially increasing LDL-C compared to placebo control.

8. The method of claim 1, comprising administering to the subject about 4 g of the pharmaceutical composition daily for the period of at least about 12 weeks to effect a reduction in apolipoprotein B compared to placebo control.

9. The method of claim 1, comprising administering to the subject about 4 g of the pharmaceutical composition daily for the period of at least about 12 weeks to effect a reduction in VLDL-C compared to placebo control.

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